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TEXTBOOK OF IMMUNOLOGY

Second Edition



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Accurate indications, adverse reactions, and dosage schedules for drugs are provided in this book, but it is possible that they may change. The reader is urged to review the package information data of the manufacturers of the medications mentioned.

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Preface to the Second Edition

We have made a series of changes in order to accommodate the new information that became available during the last four years. As before, this book has been designed as an introductory textbook with a heavy emphasis on basic immunological phenomena. The major revision concerns the new information on immunoglobulin genes, the analysis of histocompatibility and the function of T cells. The latter two are probably the most difficult topics for the students, and are presented gradually, in four chapters. First, we present a general description of the histocompatibility molecules and their genes (in Chapter 6), followed by their function in transplantation and physiological interactions (in Chapters 7, 8, and 9). Lastly, in Chapter 10, we describe the function of immune response genes as originally covered.

We are grateful to our many colleagues who helped us with advice and comments. To note is that Dr. Kurt J. Bloch revised Chapter 3, and, with Dr. Susan Canelosi, a great part of Chapter 4. Dr. K. F. Austen revised the complement chapter. Dr. F. S. D. revised Chapter 15. Dr. Ronald N. Germain wrote parts of Chapter 16. We also acknowledge the help, in this second edition, of Drs. Michael Dorf, Edmond J. Yunis, and Elvin Kabat. Both George F. Schreiber and F. S. Rosen read most of the chapters and provided considerable editorial and scientific matters. We are grateful to Battaglino, who drew all the new graphs of this edition. Finally, we owe a great deal to Barbara K. Gricus, who typed the whole book and also helped us with editorial matters.

Emil R. Unanue

Baruj Benacerraf

Boston, Massachusetts
November, 1983

prolongation of allograft or tumor survival by specific against the foreign tissue.

Immunizing agent, usually one that occurs many times on the same molecule (Hapten, Determinant, and Antigen).

Fragment of an antibody molecule consisting of two light chains and part of one heavy chain. Fab fragments combine with antigen at the antigen-binding site.

Fragment of an antibody molecule consisting of two heavy chains and two combining sites but lacking the Fc region.

Fragment of an antibody molecule consisting of two heavy chains and no combining sites for antigen. Fc has sites for activation of complement and for the binding of immunoglobulins to macrophages, lymphocytes, and mast cells. It is responsible for many biological functions of antibodies.

Protein on the surface of most lymphocytes and phagocytes that binds to the Fc portion of immunoglobulins of the IgG class.

Immunoreactive region in lymphoid tissue, in the supercortical region of lymph nodes, containing mostly B cells.

Antibodies with slow electrophoretic mobility in the gel, includes most of the immunoglobulin molecules. This term is sometimes used to refer to all immunoglobulins of various classes.

Antibodies of a large number of serum proteins distinct from immunoglobulins, insoluble at high salt concentrations.

Reaction (GVH): the pathological reactions caused by the interaction of immunocompetent T lymphocytes to an incompetent host is unable to reject the T lymphocytes and becomes a GVH reaction.

Immunocompatibility complex in the mouse.

Loci in the MHC coding for Class I histocompatibility antigens are responsible for the rapid rejection of allografts that serve as targets for T cell cytotoxicity in physiological reactions.

Hapten: chemically defined determinant that, when conjugated to an immunogenic carrier, stimulates the synthesis of specific antibody. It is capable of binding to antibody but cannot by itself stimulate an immune response.

Heavy Chain: the higher molecular weight polypeptide chain in an immunoglobulin molecule and the one determining the class of the immunoglobulin.

Helper Cells: a class of T cells that are necessary to help B cells produce antibody to thymus-dependent antigens.

Heterologous: originating from a different individual or a different inbred line; sometimes applied to a different carrier molecule (see Isologous).

Histocompatibility Antigens: cell surface antigens characteristic of an individual or an inbred line that regulate the interactions of T lymphocytes and also stimulate the rejection of tissue allografts.

HLA: the major histocompatibility complex in man.

HLA-A, HLA-B, HLA-C: three distinct genetic loci in the MHC of man coding for Class I major histocompatibility antigens.

HLA-D: a region of the MHC of man coding for Class II antigens expressed primarily on B cells and macrophages.

Humoral Immunity: immune phenomena involving the production of specific antibody.

Hypersensitivity: a poor term yet widely used, usually applied to those immune phenomena that are damaging in some way to the host animal.

Hypervariable Region: defined portions of the variable region of either heavy or light immunoglobulin chains having extreme variability in amino acid sequence in different molecules. The antibody-combining site is made of the hypervariable regions.

Ia Antigens: histocompatibility antigens found on B cells, macrophages, T cells, and Langerhans cells of the skin. They are coded for in the I region of the MHC. The Ia antigens are the Class II molecules.

Incomplete Freund's Adjuvant (ICFA): Freund's adjuvant without mycobacteria.

